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INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference GRM: FP 5793	FOR FURTHER See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416).							
International application No.	International filing date	(day/month/year)	Priority Date (day/month/year)					
PCT/AU 98/00115	CT/AU 98/00115 24 February 1999 24 February 1997							
International Patent Classification (IPC) or national classification and IPC								
Int. Cl. ⁶ C22C 21/02, 21/04								
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 This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36. 								
This REPORT consists of a total of 4 sheets, including this cover sheet.								
This report is also accompanied by ANNEXES, i.e., sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).								
•	These annexes consist of a total of sheet(s).							
3. This report contains indications relating	3. This report contains indications relating to the following items:							
I X Basis of the repor	;							
II Priority			·					
III Non-establishmen	Non-establishment of opinion with regard to novelty, inventive step and industrial applicability							
1V Lack of unity of i								
VI Certain document	s cited							
VII Certain defects in	the international applicat	tion						
VIII Certain observation	ns on the international a	pplication						
Date of submission of the demand 18 September 1998		ite of completion of January 1999	the report					
Name and mailing address of the IPEA/A AUSTRALIAN PATENT OFFICE PO BOX 200 WODEN ACT 2606 AUSTRALIA Facsimile No. (02) 6285 3929	L	J. MENZ	23. Weller 47.					

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No.

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1.	E	Basis of the report	
1.	With	regard to the elemen	its of the international application:*
	X	the international app	plication as originally filed.
		the description,	pages, as originally filed,
			pages , filed with the demand,
			pages , filed with the letter of .
		the claims,	pages , as originally filed,
	<u> </u>		pages, as amended (together with any statement) under Article 19,
			pages, filed with the demand,
			pages, filed with the letter of.
		the drawings,	pages , as originally filed,
		•	pages, filed with the demand,
			pages , filed with the letter of .
		the sequence listing	part of the description:
	1		pages , as originally filed
			pages , filed with the demand
			pages , filed with the letter of
2.	which	n the international ap	ge, all the elements marked above were available or furnished to this Authority in the language in plication was filed, unless otherwise indicated under this item. lable or furnished to this Authority in the following language which is:
		the language of a tr	ansiation furnished for the purposes of international search (under Rule 23.1(b)).
			plication of the international application (under Rule 48.3(b)).
		the language of the and/or 55.3).	translation furnished for the purposes of international preliminary examination (under Rules 55.2
3.		regard to any nucleo	otide and/or amino acid sequence disclosed in the international application, was on the basis of the
			ernational application in written form.
:			the international application in computer readable form.
	H	_	ently to this Authority in written form.
	7	•	ently to this Authority in computer readable form.
		The statement that	the subsequently furnished written sequence listing does not go beyond the disclosure in the
		The statement that	cation as filed has been furnished. the information recorded in computer readable form is identical to the written sequence listing has
		been furnished	
4.			ave resulted in the cancellation of:
		the descript	
		the claims,	Nos.
		the drawing	
5.		to go beyond the d	en established as if (some of) the amendments had not been made, since they have been considered isclosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)).**
•	Repla	cement sheets which h	ave been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this
	repor	t as "originally filed" i	and are not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17).

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No.
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V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

l	and explanations supporting s	uch statement	
	1. Statement		
	Novelty (N)	Claims 1-14 Claims	YES NO
	Inventive step (IS)	Claims 1-14 .	YES NO
	Industrial applicability (IA)	Claims 1-14 Claims	YES NO _

2. Citations and explanations (Rule 70.7)

tions

- (a) AU 41041/78
- (b) AU 35111/78
- (c) AU 86630/75

Explanations

The citations do not disclose or suggest A1-Si-Mg alloys with a microstructure including a primary aluminium containing matrix and one or more iron-containing phases dispersed in the matrix and wherein the sole or predominant iron-containing phase is β phase that formed as a transition product of π phase.

Therefore claims 1 to 14 are novel and have inventive step.

The invention, as claimed, is industrially applicable.

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VIII	Certain observations on the international application
VTII_	Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:

The broad statements of invention and the corresponding claims 1 and 5 detail an alloy composition where Fe is present in an amount up to 0.05%. Therefore the alloy includes the possibility that <u>no</u> Fe is present. However, when referring to the phase structure of the alloy, the invention is characterised by iron-containing phases requiring the presence of some Fe in the alloy.

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FOUNDRY ALLOY

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The present invention relates to an improved foundry alloy and to a method of producing an improved foundry alloy. In particular, the improved foundry alloy is an aluminium-based alloy.

used for structural or safety type applications where there is a requirement for high and consistent mechanical properties. The majority of components made from aluminium foundry alloys are made from hypocutectic aluminium-silicon-magnesium alloys containing a nominal silicon level of 7% by weight (601 and 603 designations). In simple terms these alloys are a composite of hard, discontinuous silicon particles and large, brittle iron intermetallics embedded in a ductile aluminium matrix.

There are three registered Australian compositions for strontium-modified aluminium - 7% silicon alloys. These are given in Table 1. The magnesium content of the alloys covers the range 0.25 to 0.4 wt% (601 alloys) and 0.45 to 0.7 wt% (603 alloys). The addition of magnesium allows castings to be heat treated to form magnesium silicide precipitates. These harden the matrix of the alloy to obtain the desired combination of strength and ductility.

Table 1. Registered alloy composition for strontium modified 601/603 type foundry alloys

Alloy	Si	þ.	no	Mn	Mg	Zn	Ţ	Other	Other Other	A1
Code								Each	Total	
AC601	6.5-7.5	0.20	0.05	0.05	0.30-0.40	0.05	0.20	0.05	0.05 0.15	Rem
CC601	6.5-7.5	0.20	0.05	0.05	0.25-0.35	0.05	0.20	0.05	0.05 0.15	Rem
AC603	6.5-7.5	0.15	0.05	0.03	0.45-0.7	0.05	0.20	0.05 0.15	0.15	Rem

Compositions indicate a maximum unless a range is given ♠ Compositions in weight percent.

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The main impurity found in these alloys is iron.

The iron solidifies from the eutectic liquid into a number of brittle phases.

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The two major iron-containing phases found in these alloys are the π phase (Al_Si_6Mg_3Fe) which is the predominant phase formed in high Mg content alloys and the β phase (Al_SiFe) which forms in low magnesium content alloys. The π phase forms into a script morphology while the β phase is less voluminous and forms into acicular plates. Both phases are detrimental to mechanical properties. High Mg contents (ie greater than 0.6 wt% Mg) are desirable to provide higher strength, but the presence of π phase at high Mg contents causes the ductility of the alloys to unfavourably decrease.

Conventional theories on the micro-mechanics of failure of premodified 601 and 603 alloys state that the iron rich intermetallic phases are critical in determining the fracture toughness as the silicon particles are small and round. Increases in the magnesium content of these alloys increase the amount of the π phase, which has a negative impact on the ductile properties of the alloys.

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Further, as some magnesium is contained in the π phase, the maximum volume fraction of magnesium silicide precipitates cannot be obtained. Thus, the alloys do not achieve the maximum possible strength consistent with their magnesium content.

Also, as the magnesium content of an alloy increases the magnesium content of the π phase may change leading to even greater volume fractions of the phase for a given Fe content.

It is thus concluded that the overall quality of

an alloy, as given by the quality index, decreases as increasing volume fractions of the π phase forms at increased magnesium contents. The quality index is given by the formula:

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Q.I. = UTS + 150 $log_{10} E$

where:

Q.I. = Quality Index (Mpa)

UTS = Ultimate Tensile Strength (Mpa)

E = Elongation at Fracture (%)

Attempts have been made to eliminate the π phase and thus remove its detrimental impact on mechanical properties.

By way of example, beryllium additions can be used to precipitate the iron impurity as part of the BeSiFe₂Al₈ phase. This beryllium-containing phase forms in preference to the π phase, leading to alloys with improved mechanical properties. Unfortunately, there are serious health hazards associated with using beryllium. Consequently, beryllium modification is not widely practised and the deleterious effect of the π phase on alloy quality remains.

Other attempts to modify the iron-containing phases, for example by using Mn, have been tried in secondary alloys but have not been used in primary alloys.

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It is an object of the present invention to provide an improved foundry alloy.

In accordance with the present invention this object is achieved by an alloy having a microstructure in which β phase that forms during heat treatment as a transformation product of π phase is the sole or

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predominant iron-containing phase. The reduction in π phase results in an improvement in ductility. the β phase that forms as the transformation product has a fine structure that improves ductility. Further, the reduction in π phase means that there are higher levels of Mg in solution which are available for precipitation during ageing to improve the strength of the alloy.

In a first aspect, the present invention provides an alloy which comprises: 10

> 6.5 - 7.5 wt% Si up to 0.20 wt% Fe Cu up to 0.05 wt% up to 0.05 wt% Mn 0.35 to 0.50 wt%

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up to 0.05 wt% z_n : Ti up to 0.20 wt% :

Mg

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Balance: Al and other components, the other 20 components comprise a total of not more than 0.15 wt% and any single component of the other components does not exceed 0.05 wt%, the alloy having a microstructure which includes a primary aluminium-containing matrix and one or more iron-containing phases dispersed in the matrix, and 25 wherein the sole or predominant iron-containing phase is β phase that formed as a transformation product of π phase.

It is preferred that the dendrite arm spacing of 3.0 the matrix be 10-45µm.

Where there is more than one iron-containing phase, preferably the iron-containing phases also include π phase.

Preferably, the iron-containing phases include π phase in an amount up to 30 vol% of the iron-containing

phases. The amount of π phase may be higher if the Mg content is in the upper end of the range.

The Mg content of the alloy is preferably 0.40-0.45 wt%. Within this Mg range, the alloy is a variant of the 601/603 type foundry alloy. It has been realised by the applicant that close control of the magnesium content to be between 0.40 and 0.45 wt% can lead to an increase in alloy quality and improved mechanical properties. 10 particular, when the magnesium content is controlled to be between 0.40 and 0.45 wt% the variation in alloy quality for a small change in magnesium level is minimal. the consistency in the mechanical properties of the alloy is maximised.

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The present invention also provides a method for manufacturing an alloy article.

In a second aspect, the present invention provides a method for manufacturing an alloy article which 20 comprises:

> providing a melt having a composition of: (a)

Si 6.5 - 7.5 wt% up to 0.20 wt% Fe

up to 0.05 wt% Cu

up to 0.05 wt% Mn

0.35 to 0.50 wt% Mg

up to 0.05 wt%

Ti up to 0.20 wt%

 $\mathbf{Z}\mathbf{n}$

Balance : Al and other components, the other components comprising a total of not more than 0.15wt% and any single component of the other components not exceeding 0.05 wt%,

(b) casting said melt and solidifying a casting WO 98/38347 PCT/AU98/00115

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at a cooling rate that produces a microstructure of an aluminium-containing matrix and π and β iron-containing phases dispersed in the matrix;

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(c) solution heat treating the casting to at least partially transform π phase to β phase; and

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(d) quenching the casting to form the alloy article.

It is preferred that the cooling rate be sufficient to produce a dendrite arm spacing in the matrix in the casting of $10-45\mu m$.

Preferably, the sole or predominant ironcontaining phase in the alloy article is β phase.

20 Where there is more than one iron-containing phase in the alloy article, preferably the iron-containing phases also include π phase. More preferably, the iron-containing phases in the alloy article include π phase in an amount of up to 30 vol% of the iron-containing phases.

25 Higher levels of π phase may be present if the Mg content

is at the upper end of the above range.

It is preferred that the step of solidifying the casting produces iron-containing phases that include a substantial proportion of the π phase and the subsequent solution heat treatment step is effective to convert at least some and preferably a majority of the π phase to β phase to give a microstructure in the alloy article that includes iron-containing phases which are predominantly β phase.

The melt prior to casting may be at a temperature

above the liquidus temperature of the alloy, with the melt having sufficient superheat to fill the mould, that is at a temperature of 680-720°C.

The solution treatment of the casting may be carried out at any suitable temperature and for any suitable time to achieve a desired level of transformation of π phase β phase. In any given situation, the selection of the parameters of temperature and time will depend on variables, such as the concentrations of magnesium and other elements in the casting. By way of example, the applicant has found that for castings having a Mg concentration of 0.5 wt%, solution treatment at 540°C for 2 or more hours produced desired levels of transformation of π to π phase

After the solution heat treatment, the casting is preferably quenched, more preferably quenched in hot water, such as hot water having a temperature of 70-80°C.

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After quenching, the alloy article is cooled to room temperature and optionally subjected to an ageing heat treatment.

The ageing heat treatment may include heating the alloy article to a temperature of 140-170°C and holding at that temperature for 1-10 hours. After the ageing heat treatment, the alloy article may be air cooled to room temperature.

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Results to support the present invention are given in Figure 1, in which plots of typical response surfaces derived from experimentally determined quality index data are shown. The three surfaces correspond to alloys that were cast at different solidification rates and thereafter solution treated and aged. Solidification rate is commonly measured by the as-cast dendrite cell size or

secondary dendrite arm spacing (DAS) but other methods exist. The results here use secondary dendrite arm spacing to indicate solidification rate, with a small dendrite arm spacing corresponding to a high solidification rate.

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It can be seen from Figure 1 that:

- (i) at the high solidification rate (≈20μm DAS)
 the alloy quality peaks at a magnesium level
 of 0.45-0.50 weight percent;
 - (ii) at the intermediate solidification rate ($\approx 40 \mu m$ DAS) the quality peaks at a magnesium level of 0.35-0.40 weight percent; and

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(iii) at the low solidification rate (≈60µm DAS) the quality maximum occurs at a magnesium level of 0.25-0.30 weight percent.

Further, it can be seen from Figure 1 that the magnesium level for the peak quality is independent of the iron level for the iron levels examined. Also, the rate of change of the response surfaces with magnesium is least near the peak in quality index. This means that the alloys at the peak are less sensitive to changes in magnesium than other alloys. The peak quality from Figure 1 corresponds well with microstructural evidence for small amounts of π phase in the alloy. By increasing the magnesium content of the alloy, it can be seen that in some circumstances improved quality results.

It should be noted that the present invention works best with those casting designs or casting methods which produce high solidification rates ($\leq 45 \mu m$ DAS), such as permanent mould, mould chill methods with sand, and squeeze casting. Indeed, the trend in the automotive industry is to move away from thick section, low

solidification rate (high DAS) castings towards lightweight castings with thinner sections and higher solidification rates (low DAS).

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The common belief prior to the present invention was that low magnesium levels produce high quality castings. The results shown here confirm this to be true at low solidification rates (Figure 1c). However, at higher solidification rates, the magnesium contents covered by this invention show, surprisingly, improved alloy quality and therefore improved mechanical properties.

Figures 2(a) to 2(c) are photomicrographs of hypoeutectic alloys having a Si concentration of 7 wt% and various Mg concentrations which were cast at the same solidification rate (60µm DAS), solution treated, and aged. Figure 2(d) is a photomicrograph of the as-cast alloy of Figure 2(c), ie before heat treatment.

In Figure 2(a), the Mg content of the alloy is higher than the Mg content of the alloy of the present invention. The main phases shown in Figure 2(a) are spheroidal silicon-containing phase and the iron-containing π phase.

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Figure 2(b) shows the microstructure of an alloy containing less Mg than the alloy of the present invention. The phases present include spheroidal silicon-containing phase and iron-containing β phase. The β phase is present as structures of high aspect ratio dispersed throughout the matrix.

Figure 2(c) shows the microstructure of an alloy of the present invention. The phases include spheroidal silicon-containing phases, a small amount of π phase and β phase. The β phase is present as structures of high aspect ratio clumped together. This is consistent with the β

phase being formed by transformation of π phase during heat treatment.

Figure 2(d) shows that prior to heat treatment the as-cast alloy of Figure 2(c) had regions of π phase. As is evident from Figure 2(c) these π phase regions were largely transformed to β phase during heat treatment.

The drive for alloys with improved mechanical properties stems from the major restraint that mechanical 10 properties place on the design of the casting, or even if a cast alloy can be used to manufacture a certain component. The thickness of critical sections needs to be sufficiently large that the cast component can operate without failure. 15 Mechanical properties of the alloys therefore limit the minimum weight of a cast component. Further, the thickness of sections of a casting will determine the time required for the casting to solidify. For certain casting methods, such as low pressure die casting, the production rate is often determined by the solidification rate as the casting 20 machine is tied up until the casting has fully solidified. Finally, the solution treatment, quench rate and ageing treatment of a cast component may be tailored to its design so as not to induce unnecessarily high residual stresses. 25 High residual stresses can cause distortion of the component requiring additional machining. The mechanical properties of the base alloy therefore affect all stages of

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The present invention therefore has the following more specific applications:

manufacturing from design, to casting the component, heat treatment, machining, final weight and production rate.

(i) New markets for aluminium-7% silicon foundry

alloys. Cast alloys generally have inferior
mechanical properties but lower
manufacturing costs compared to similar

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components made from wrought alloys. The high mechanical property requirements of some components necessitates the use of wrought alloys. The achievement of alloys of the present invention which have higher and more consistent mechanical properties than conventional alloys may allow the use of the alloy of the present invention to replace wrought alloys, or other cast alloys, for some components.

- (ii) Cast components with thinner sections and lower total weight. The improved and more consistent mechanical properties of the alloy of the present invention allows components with thinner sections to be designed and cast. Despite their thinner sections, these components can still operate without failure and will have a lower total weight.
- (iii) Cast components with an improved production rate. Castings with thinner sections may require less time to solidify. Production equipment will then be tied up for less time waiting for a component to solidify. The production rate is thus increased.
- (iv) Cast components with refined iron and silicon intermetallic phases. The solidification time of a casting strongly determines the coarseness of the microstructure. Components with thinner sections and therefore higher solidification rates (and lower solidification times) will have a more refined microstructure. This refining of the microstructure will provide

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additional improvements to the mechanical properties of a casting, independent of the use of a superior alloy.

- (v) Cast components with reduced heat treatment time. Castings with thinner sections require less time to homogenise. Further, the time required for the casting to reach the solution treatment temperature or ageing temperature will be less. This also benefits the production rate of components.
 - (vi) Cast components with increased quench rate. Thinner castings may quench more rapidly. This may lead to improved mechanical properties as it suppresses the formation of magnesium-silicide precipitates during cooling. These improved properties are independent of any refinement of the microstructure or the use of a superior alloy.

It will be appreciated that the invention described herein is susceptible to variation and modifications other than those specifically described. It is to be understood that the invention encompasses all such variations and modifications that fall within its spirit and scope.

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CLAIMS:

An alloy which comprises:

5 Si : 6.5 - 7.5 wt%

Fe : up to 0.20 wt%

Cu : up to 0.05 wt%

Mn : up to 0.05 wt%

Mg : 0.35 to 0.50 wt%

10 Zn : up to 0.05 wt%

Ti : up to 0.20 wt%

Balance: Al and other components, the other components comprise a total of not more than 0.15 wt% and any single component of the other components does not exceed 0.05 wt%, the alloy having a microstructure which includes a primary aluminium-containing matrix and one or more iron-containing phases dispersed in the matrix, and wherein the sole or predominant iron-containing phase is β phase that formed as a transformation product of π phase.

- 2. The alloy defined in claim 1, wherein when the alloy includes more than one iron-containing phase, the iron-containing phases also include π phase.
 - 3. The alloy defined in claim 2, wherein the π phase is up to 30 vol% of the iron-containing phases.
 - 4. The alloy defined in any one of the preceding claims, wherein the Mg content of the alloy is 0.40-0.45 wt%.
- 35 5. A method for manufacturing an alloy article which comprises:

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(a)	prov	riding	g a melt having a composition o
	si	:	6.5 - 7.5 wt%
	Fe	:	up to 0.20 wt%
	Cu	:	up to 0.05 wt%
	Mn	:	up to 0.05 wt%
	Mg	:	0.35 to 0.50 wt%
	Zn	:	up to 0.05 wt%
	Ti	:	up to 0.20 wt%

Balance: Al and other components, the other components comprising a total of not more than 0.15wt% and any single component of the other components not exceeding 0.05 wt%,

- (b) casting said melt and solidifying a casting at a cooling rate that produces a microstructure of an aluminium-containing matrix and π and β iron-containing phases dispersed in the matrix;
- (c) solution heat treating the casting to at least partially transform π phase to β phase; and
- 25 (d) quenching the casting to form the alloy article.
- 6. The method defined in claim 5, wherein the cooling rate is sufficient to produce a dendrite arm
 30 spacing in the matrix of between 10 and 45μm.
 - 7. The method defined in claim 5 or claim 6, wherein the sole or predominant iron-containing phase in the alloy article is β phase.
 - 8. The method defined in claim 5, wherein when the alloy includes more than one iron-containing phase in

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the alloy article, the iron-containing phases also include π phase.

- 9. The method defined in claim 8, wherein the π 5 phase is up to 30 vol% of the iron-containing phases.
 - 10. The method defined in claim 5 or claim 6, wherein the step of solidifying the casting produces iron-containing phases that include a substantial proportion of π phase and the subsequent solution heat treatment step is effective to convert a majority of the π phase to β phase to give a microstructure in the alloy article that includes iron-containing phases which are predominantly β phase.
- 11. The method defined in any one of claims 5 to 10, wherein prior to casting the melt is at a temperature above the liquidus temperature of the alloy.
- 12. The method defined in any one of claims 5 to 20 11, wherein the quenching step is in hot water having a temperature of 70-80°C.
- 13. The method defined in any one of claims 5 to 12, further includes an ageing heat treatment of the alloy 25 article.
 - 14. The method defined in claim 13, wherein the ageing heat treatment includes heating the alloy article to a temperature of 140-170°C, holding the alloy article at that temperature for 1-10 hours, and air cooling the alloy article to room temperature.

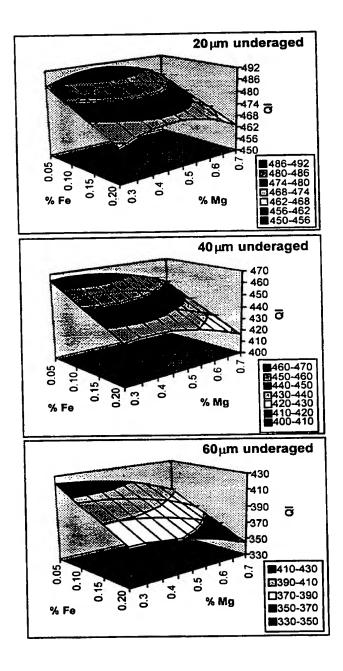
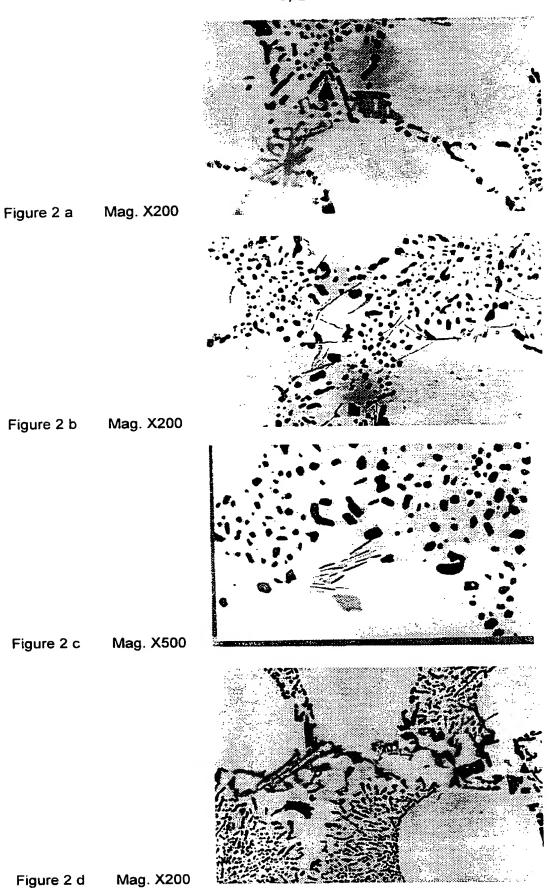
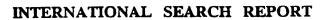


Figure 1





International Application No. PCT/AU 98/00115

A.	CLASSIFICATION OF SUBJECT MATTER								
Int Cl ⁶ :	C22C 21/02, 21/04								
According to International Patent Classification (IPC) or to both national classification and IPC									
	FIELDS SEARCHED								
	Minimum documentation searched (classification system followed by classification symbols) C22C 21/02, C22C 21/04								
_	searched other than minimum documentation to the ext as above	ent that such documents are included in t	he fields searched						
	base consulted during the international search (name of se - alloy search (A1 and 0.35<=mg<0.5 a # or Zone #)	data base and, where practicable, search and 6.5<=Si<=7.5 and Fe<=0.2) an							
C.	DOCUMENTS CONSIDERED TO BE RELEVANT								
Category*	Citation of document, with indication, where appropriate, of the relevant passages Relevant to claim No.								
A	AU,A, 41041/78 (AU,B, 515605) (ALUMINIUM COMPANY OF AMERICA) 1 May 1980 In General								
A	AU,A, 35111/78 (MESSIERS S A) 18 October 1979 In General								
A	AU,A, 86630/75 (AU,B, 507432) (ALCAN RESEARCH AND DEVELOPMENT) 19 May 1977 In General								
Further documents are listed in the continuation of Box C									
"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention cannot be considered novel or cannot be considered to invention cannot be considered novel or cannot be considered to invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means document published prior to the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention cannot be considered novel or cannot be considered to involve an inventive step when the document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art document member of the same patent family									
	ual completion of the international search	Date of mailing of the international sear - 8 MAY	- /						
0.1417/4000									



Information on patent family members

International Application No. PCT/AU 98/00115

This Annex lists the known "A" publication level patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent Do	cument Cited in Search Report			Patent	Family Member		
AU	41041/78	CA	1084661	DE	2848653	FR	2415002
		GB	2007547	JP .	54099762	US	4146163
AU	35111/78	СН	624716	FR	2388892		
AU	86630/75	AT	8708/75	BE	835582	BR	7507572
		CA	1041880	CH	60 64 60	DE	255129 5
		DK	5124/75	ES	442619	FR	2291284
		GB	1529305	HK	411/79	IN	149783
		Π	1049091	JР	51073913	MY	2/83
		NL	7513351	NO	753833	NO	794344
		SE	7512813	US	4126486	YU	2831/75
		ZA	7506999				

END OF ANNEX

28 Rec'd PCT/PTO PAG TUG 1999

REQUEST

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HPCT/PTOPAGTAUG1999	Г	For	receiv	ing Office use only			
Mon teligies les	lnte	emational Applicatio	n No.				
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REQUEST	Int	ernational Filing Dat	le				
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The undersigned requests that the present international application be processed	N	Name of receiving Office and "PCT International Application"					
according to the Patent Cooperation Treaty.		Applicant's or agent's file reference (if desired):(12 cliaracters maximum) GRM: FP5793					
x No. I TITLE OF INVENTION							
MPROVED FOUNDRY ALLOY							
x No. II APPLICANT							
ame and address; (Family name followed by given name: for a le le address must include postal code and name of country. The country ox is the applicant's State (i.e. country) of residence if no State of r	gal entitions of the residence	y, full official designation e address indicated in the e is indicated below.)		This person is also inventor.			
AST CENTRE PTY LTD			T	elephone No.			
:/- DEPARTMENT OF MINING FTALLURGICAL ENGINEERING			F	Facsimile No.			
THE UNIVERSITY OF QUEENSLAND ST LUCIA, QUEENSLAND 4072			T	Teleprinter No.			
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his person is applicant all designated all designated the University of the purposes of:	nited Sta	States except	of A	merica only the Supplemental Box			
30x No. III FURTHER APPLICANT(S) AND/OR (F	URTH	ER) INVENTOR(S	»)				
Name and address: (Family name followed by given name: for a he address must include postal code and name of country. The cou Box is the applicant's State (i.e. country) of residence if no State of	legal en uniry of residen	icy, full official designations and cated in the cardinated in the cated in the cated below.)	this	This person is: applicant only			
BARRESI, JOSEPH GIOVANNI 17 MARY BYRANT COURT				x applicant and inventor			
MILL PARK, VICTORIA 3082 AUSTRALIA				inventor only (If this check-bax is marked, do not fill in below.)			
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Further applicants and/or (further) inventors are indi	icated o	n a continuation she	et.				
Box No. IV AGENT OR COMMON REPRESENTA							
The person identified below is hereby/has been appointed of the applicant(s) before the competent International Authority				gent common representative			
Name and address: (Family name followed by given name; for The address must include postal code and	raiegali d name i	of country.)		(61 03)9243 8300			
GRIFFITH HACK 509 ST KILDA ROAD MELBOURNE, VICTORIA 3004				Facsimile No. (61 03)9243 8333			
AUSTRALIA	٠			Teleprinter No.			

indicate a special address to which correspondence should be sent. Form PCT/RO/101 (first sheet) (January 1997; reprint July 1997)

See Notes to the request form

2 Sheet No. FURTHER APPLICANTS AND/OR (FURTHER) INVENTORS Continuation of Box No. III If none of the following sub-boxes is used, this sheet is not to be included in the request. Name and address: (Family name followed by given name: for a legal entire, full official designation. The address must include postal code and name of country. The country of the address indicated in this Box is the applicant's State (i.e. country) of residence if no State of residence is indicated below.) This person is: applicant only COUPER, MALCOLM JAMES 49 ORONSAY CRESCENT applicant and inventor X DIAMOND CREEK, VICTORIA 3089 inventor only (If this check-box AUSTRALIA is marked, do not fill in below.) State (i.e. country) of residence: State (i.e. country) of nationality: AUSTRALIA <u>AUSTRALI</u>A the States indicated in the Supplemental Box the United States all designated States except the United States of America all designated States This person is applicant of America only for the purposes of: Name and address: (Family name followed by given name; for a legal entiry full official designation. The address must include postul code and name of country. The country of the address indicated in this Box is the applicant's State (i.e. country) of residence if no State of residence is indicated below.) This person is: applicant only ST JOHN, DAVID HENRY applicant and inventor 26 GLADSTONE STREET INDOOROOPILLY, QUEENSLAND 4068 inventor only (If this check-box AUSTRALIA is marked, do not fill in below.) State (i.e. country) of residence: State (i.e. country) of nationality: AUSTRALIA AUSTRALIA the States indicated in the Supplemental Box the United States of America only all designated States except the United States of America Х all designated This person is applicant for the purposes of: Name and address: (Family name followed by given name: for a legal entiry, full official designation. The address must include postal code and name of country. The country of the address indicated in this Bux is the applicant's State (i.e. country) of residence if no State of residence is indicated below.) This person is: applicant only EDWARDS, GEOFFREY ALAN 18 HIGHVIEW TERRACE applicant and inventor DAISY HILL, QUEENSLAND 4127 inventor only (If this check-box is marked, do not fill in below.) AUSTRALIA State (i.e. country) of residence: State (i.e. county) of nationality: the States indicated in the Supplemental Box the United States of America only all designated States except the United States of America all designated States This person is applicant for the purposes of: Name and address: (Family name followed by given name: for a legal entity, full official designation. The address must include postal code and name of country. The country of the address indicated in this Box is the applicant's State (i.e. country) of residence if no State of residence is indicated below.) This person is: applicant only WANG, HAO applicant and inventor 14/29 VILLA STREET ANNERLEY, QUEENSLAND 4103 inventor only (If this check-box **AUSTRALIA** is marked, du not fill in below.) State (i.e. country) of residence: State (i.e. country) of nationality: AUSTRALIA CHINA the States indicated in the Supplemental Box the United States of America only all designated States except the United States of America all designated This person is applicant for the purposes of: Further applicants and/or (further) inventors are indicated on another continuation sheet.

Sheet No. . . 3

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Box No	۷.ن	DESIGNATION OF STATES			" the short horses at least one must be marked):			
The fo	llowir	ng designations are hereby made under Rule 4.9(a) (mark	the ap	plicable creek-boxes, at teast one			
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under the PCT except the designation(s) of
The applicant declares that those additional designations are subject to confirmation and that any designation which is not confirmed before the expiration of 15 months from the priority date is to be regarded as withdrawn by the applicant at the expiration of that time limit. (Confirmation of a designation consists of the filling of a notice specifying that designation and the payment of the designation and confirmation fees. Confirmation must reach the receiving Office within the 15-month time limit.)

Sheet No. ...4

No. VI PRIORITY C		ther priority claims are indicated i	n the Supplemental Box
e priority of the following es	arlier application(s) is hereby claimed	:	Office of filing
Country in which, or for which, the application was filed)	Filing Date (day/month/year)	Application No.	(unly for regional or international application)
m(I) AUSTRALIA	24.02.97 24 FEBRUARY 1997	PO5268	
m (2)			
:m (3)			
The receiving Office is Bureau a certified copy	hereby requested to prepare and trans of the earlier application(s) identified	amit to the International diabove as item(s):	purposas of the present marmamata
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AUSTRALIA	26 MAY 1997	97/55	8
Box No. VIII CHECK LE	ST		
This international application following number of shifted following number of shifted following number of shifted following in the following number of shifted number of shifted in the following number of shifted number of shifted number	sheets	of attorney of general of attorney thent explaining f signature ty document(s) fied in Box No. VI	e calculation sheet parate indications concerning posited microorganisms ucleotide and/or amino acid quence listing (diskette) ther (specify):
Figure No. 1 of	the drawings (if any) should accompa	any the abstract when it is publish	ed
	E OF APPLICANT OR AGENT		
Next to each signature, indicate the	name of the person rigning and the capacity Authorised signal CAST CENTRE PTY	EXECUTIVE DIRE	_
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Date of timely receipt of corrections under PCT.			not receive
5. International Searching	Authority ISA /	6. Transmittal of search countil search fee is paid	opy delayed
specified by the applica	nt:		

Sheet No. ... 5...

Supplemental Box If the Supplemental Box is not used, this sheet need not be included in the request.

Use this box in the following cases:

 If, in any of the Boxes, the space is insufficient to furnish all the information:

in particular:

- (i) if more than two persons are involved as applicants and/or inventors and no "continuation sheet" is available:
- (ii) if, in Box No. II or in any of the sub-boxes of Box No. III, the indication "the States indicated in the Supplemental Box" is checked:
- (iii) if, in Box No. If or in any of the sub-boxes of Box No. III, the inventor or the inventor/applicant is not inventor for the purposes of all designated States or for the purposes of the United States of America:
- (iv) if, in addition to the agent(s) indicated in Box No. IV, there are further agents:
- (v) if, in Box No. V. the name of any State (or OAPI) is accompanied by the indication "patent of addition," or "certificate of addition," or if, in Box No. V. the name of the United States of America is accompanied by an indication "Continuation" or "Continuation in-part".
- (vi) if there are more than three earlier applications whose priority is claimed:

2. If the applicant claims, in respect of any designated Office, the benefits of provisions of the national law concerning non-prejudicial disclosures or exceptions to lack of novelty:

in such case, write "Continuation of Box No. ..." [indicate the number of the Box] and furnish the information in the same manner as required according to the captions of the Box in which the space was insufficient;

in such case, write "Continuation of Box No. III" and indicate for each additional person the same type of information as required in Box No. III. The country of the address indicated in this Box is the applicant's State (i.e. country) of residence if no State of residence is indicated below:

in such case, write "Continuation of Box No. II" or "Continuation of Box No. III" or "Continuation of Boxes No. II and No. III" (as the case may No. III" or "Continuation of Boxes No. II and No. III" (as the case may be), indicate the name of the applicant(s) involved and, next to (each) such name, the State(s) (and/or, where applicable, ARIPO. Eurasian, European or OAPI patent) for the purposes of which the named person is applicant:

in such case, write "Continuation of Box No. II" or "Continuation of Box No. III" or "Continuation of Boxes No. II and No. III" (as the case may No. III" or "Continuation of Boxes No. II and No. III" (as the case may be), indicate the name of the inventor(s) and, next to (each) such name, the State(s) (and/or, where applicable, ARIPO, Eurasian, European or OAPI parent) for the purposes of which the named person is inventor:

in such case, write "Continuation of Box No. IV" and indicate for each further agent the same type of information as required in Box No. IV:

in such case, write "Continuation of Box No. V" and the name of each State involved (or OAPI), and after the name of each such State (or OAPI), the number of the parent title or parent application and the date of grant of the parent title or filing of the parent application:

in such case, write "Continuation of Box No. VI" and indicate for each additional earlier application the same type of information as required in Box No. VI.

in such case, write "Statement Concerning Non-Prejudicial Disclosures or Exceptions to Lack of Novelty" and furnish that statement below.

Continuation of Box No. IX (Signatures)

V JOSEPH GEOVANNI BARRESI

MALCOLM JAMES COUPER

DAVID HENRY ST JOHN

MULLIUM:

GEOFFREY ALAN EDWARDS

HAO WANG

WiFo

2 9 JAN 1999

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference GRM: FP 5793	FOR FURTHER ACTION		f Transmittal of International Preliminary ort (Form PCT/IPEA/416).
International application No.	International filing date	e (day/month/year)	Priority Date (day/month/year)
PCT/AU 98/00115	24 February 1999		24 February 1997
International Patent Classification (IPC)	or national classification	n and IPC	
Int. Cl. ⁶ C22C 21/02, 21/04			
Applicant (1) CAST CENTRE PTY LTD (2) BARRESI, Joseph Giovanni (et al)			
This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.			
2. This REPORT consists of a to	tal of 4 sheets, includ	ling this cover sheet.	
This report is also accompanied by ANNEXES, i.e., sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).			
These annexes consist of a total			
3. This report contains indications relati	ng to the following item	s:	
I X Basis of the repor	I X Basis of the report		
II Priority	II Priority		
III Non-establishmen	III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability		
IV Lack of unity of i	IV Lack of unity of invention		
V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement			
VI Certain documen	nts cited		
VII Certain defects in	n the international application		
VIII Certain observati	VIII Certain observations on the international application		
Date of submission of the demand 18 September 1998		Date of completion of 2 January 1999	f the report
Name and mailing address of the IPEA/AU AUSTRALIAN PATENT OFFICE PO BOX 200 WODEN ACT 2606 AUSTRALIA		Authorized Officer 23. Weller L.J. MENZ	
Facsimile No. (02) 6285 3929		Telephone No. (02) 6283 2347	

I.	I	Basis of the report	
1.	With	regard to the elements of	he international application:*
	X	the international applicati	on as originally filed.
		the description, page page	
		page	s, as originally filed, s, filed with the demand, s, filed with the letter of .
		the sequence listing part of pages pages	, as originally filed , filed with the demand
2.	which	the international applicati	the elements marked above were available or furnished to this Authority in the language in on was filed, unless otherwise indicated under this item. or furnished to this Authority in the following language which is:
		the language of publication	ion furnished for the purposes of international search (under Rule 23.1(b)). on of the international application (under Rule 48.3(b)). ation furnished for the purposes of international preliminary examination (under Rules 55.2
3.		and/or 55.3). regard to any nucleotide ance listing:	nd/or amino acid sequence disclosed in the international application, was on the basis of the
		filed together with the int	onal application in written form. ernational application in computer readable form. this Authority in written form.
		furnished subsequently to The statement that the sui international application a	this Authority in computer readable form. Description of the disclosure in the disc
4.		The amendments have remarks the description, the claims, the drawings,	pages Nos. sheets/fig
5.			plished as if (some of) the amendments had not been made, since they have been considered re as filed, as indicated in the Supplemental Box (Rule 70.2(c)).**
*	report	as "originally filed" and are	n furnished to the receiving Office in response to an invitation under Article 14 are referred to in this not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17). uch amendments must be referred to under item 1 and annexed to this report

V.	Reasoned statement under Ar and explanations supporting s			ntive step or industrial applicability; citations
1.	Statement -			
	Novelty (N)	Claims Claims	1-14	YES NO
	Inventive step (IS)	Claims Claims	1-14	YES NO
	Industrial applicability (IA)	Claims Claims	1-14	YES
2.	Citations and explanations (Rule	70.7)		

Citations

- AU 41041/78 (a)
- (b) AU 35111/78
- AU 86630/75 (c)

Explanations

The citations do not disclose or suggest A1-Si-Mg alloys with a microstructure including a primary aluminium containing matrix and one or more iron-containing phases dispersed in the matrix and wherein the sole or predominant iron-containing phase is β phase that formed as a transition product of π phase.

Therefore claims 1 to 14 are novel and have inventive step.

The invention, as claimed, is industrially applicable.

VIII. Certain observations on the international application				
The following observations on the clarity of the claims, description, and drawings or on the que supported by the description, are made:	estion whether the claims are fully			
The broad statements of invention and the corresponding claims 1 and 5 detail an alloy composition where Fe is present in an amount up to 0.05%. Therefore the alloy includes the possibility that no Fe is present. However, when referring to the phase structure of the alloy, the invention is characterised by iron-containing phases requiring the presence of some Fe in the alloy.				
F *				